FILE 'HOME' ENTERED AT 16:45:23 ON 01 MAR 2005

- L1 QUE (EPSTEIN-BARR OR EPSTEIN (A) BARR) (A) VIRUS (P) ((INDUC? OR SPECIFIED) (S) (GENE OR PROTEIN) OR VIRUS-INDUC? OR VIRUS-SPECIF?)
- L2 10174 (EPSTEIN-BARR OR EPSTEIN (A) BARR) (A) VIRUS (P) ((INDUC? OR SPECIFIED) (S) (GENE OR PROTEIN) OR VIRUS-INDUC? OR VIRUS-SPECIF?)
- L3 3207 (EBV-INDUC#### OR EBV-SPECIFIED) OR EBV (A) (INDUC### OR SPECIFI ED
- L5 7182 (L1 OR L2) AND ((EPSTEIN (A) BARR)/TI OR EBV/TI OR EPSTEIN-BARR/TI)
- L8 580 L7 AND ((VIRUS-INDUCED OR VIRUS-SPECIFIED) OR (VIRUS OR EBV OR EPSTEIN (A) BARR) (3N) (INDUCED OR SPECIFIED OR REGULAT### OR ACTIVAT###))
- L9 148 L7 AND ((VIRUS-INDUCED OR VIRUS-SPECIFIED) OR (VIRUS OR EBV OR EPSTEIN (A) BARR) (3N) (INDUCED OR SPECIFIED OR REGULAT### OR ACTIVAT###)) (S) (PROTEIN OR RECEPTOR OR GENE)
- L10 124 L9 AND (EBV? OR EPSTEIN-BARR OR EPSTEIN (A) BARR) (S) CELL
- L11 47 L9 AND (VIRUS-INDUCED/TI OR VIRUS-SPECIFIED/TI OR INDUCED/TI OR SPECIFIED/TI OR REGULATED/TI)

(FILE 'HOME' ENTERED AT 16:45:23 ON 01 MAR 2005)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHOS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DISSABS, ...' ENTERED AT 16:47:31 ON 01 MAR 2005 SEA (EPSTEIN-BARR OR EPSTEIN (A) BARR) (A) VIRUS (P) ((INDUC? O

71 FILE ADISCTI

-----**---**

- 7 FILE ADISINSIGHT
- 8* FILE ADISNEWS
- 9 FILE AGRICOLA
- 1 FILE ANABSTR
- 0* FILE ANTE
- 0* FILE AQUALINE
- 1 FILE AQUASCI
- 5 FILE BIOBUSINESS
- 2* FILE BIOCOMMERCE
- 91* FILE BIOENG
- 1784 FILE BIOSIS
- 279* FILE BIOTECHABS
- 279* FILE BIOTECHDS
- 1687* FILE BIOTECHNO
 - 44 FILE CABA
- 2096 FILE CANCERLIT
- 1511 FILE CAPLUS
 - 10* FILE CEABA-VTB
 - 3* FILE CIN
 - 46 FILE CONFSCI
 - 34 FILE DDFB
- 102 FILE DDFU
- 2697 FILE DGENE
- 113 FILE DISSABS
 - 34 FILE DRUGB

```
FILE EMBASE
       2160
              FILE ESBIOBASE
       1127*
         78*
              FILE FEDRIP
          0*
              FILE FOMAD
              FILE FOREGE
          0*
          0*
              FILE FROSTI
          2*
              FILE FSTA
         65
              FILE GENBANK
          3
              FILE HEALSAFE
         87
              FILE IFIPAT
          2
              FILE IMSDRUGNEWS
              FILE IMSRESEARCH
          6
              FILE JICST-EPLUS
         84
              FILE KOSMET
          0*
       1293
              FILE LIFESCI
          0*
              FILE MEDICONF
       1607
              FILE MEDLINE
          3
              FILE NIOSHTIC
          8*
              FILE NTIS
          0*
             FILE NUTRACEUT
             FILE PASCAL
        *888
              FILE PHAR
          1
              FILE PHARMAML
          0*
              FILE PHIN
          2
         48
              FILE PROMT
              FILE PROUSDDR
          1
          1
              FILE RDISCLOSURE
       1819
              FILE SCISEARCH
        790
              FILE TOXCENTER
        829
              FILE USPATFULL
              FILE USPAT2
         55
              FILE VETU
          2
             FILE WATER
          0*
        131
              FILE WPIDS
              FILE WPINDEX
           QUE (EPSTEIN-BARR OR EPSTEIN (A) BARR) (A) VIRUS (P) ((INDUC? O
FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, LIFESCI' ENTERED AT
16:54:37 ON 01 MAR 2005
     10174 S (EPSTEIN-BARR OR EPSTEIN (A) BARR) (A) VIRUS (P) ((INDUC? OR
      3207 S (EBV-INDUC#### OR EBV-SPECIFIED) OR EBV (A) (INDUC### OR SPEC
     10174 S (L1 OR L2) AND (EPSTEIN (A) BARR (S) VIRUS)
      7182 S (L1 OR L2) AND ((EPSTEIN (A) BARR)/TI OR EBV/TI OR EPSTEIN-BA
      2270 DUP REM L5 (4912 DUPLICATES REMOVED)
      1032 S L6 AND PY<1993
      580 S L7 AND ((VIRUS-INDUCED OR VIRUS-SPECIFIED) OR (VIRUS OR EBV
      148 S L7 AND ((VIRUS-INDUCED OR VIRUS-SPECIFIED) OR (VIRUS OR EBV
      124 S L9 AND (EBV? OR EPSTEIN-BARR OR EPSTEIN (A) BARR) (S) CELL
```

47 S L9 AND (VIRUS-INDUCED/TI OR VIRUS-SPECIFIED/TI OR INDUCED/TI

177

L1

L2

L3

L4

L5 L6

L7

L8

L9 L10

L11

24

FILE DRUGU

FILE EMBAL

((VIRUS-INDUCED OR VIRUS-SPECIFIED) OR (VIRUS OR EBV OR EPSTEIN (A) BARR) (3N) (INDUCED OR SPECIFIED OR REGULAT###

ANSWER 7 OF 47 L11 MEDLINE on STN MEDLINE AN87214791 PubMed ID: 3034370 DN The prevalence of antibodies to an Epstein-Barr ТT virus-induced polypeptide (EBNA-2) in the sera of rheumatoid arthritic families. ΑU Hazelton R A; Sculley T B; Pope J H British journal of rheumatology, (1987 Jun) 26 (3) 193-6. SO Journal code: 8302415. ISSN: 0263-7103. CYENGLAND: United Kingdom DT Journal; Article; (JOURNAL ARTICLE) LA FS Abridged Index Medicus Journals; Priority Journals ΕM 198706 ED Entered STN: 19900303 Last Updated on STN: 19900303 Entered Medline: 19870629 Using the protein immunoblot technique, antibodies to an AB Epstein-Barr virus-induced 92 kD polypeptide (EBNA-2) were more frequently present in the sera of patients with rheumatoid arthritis and their consanguineous relatives when compared with a control group. No association of anti-EBNA-2 antibody with the HLA-DR antigens was observed. L11 ANSWER 8 L11 ANSWER 10 OF 47 MEDLINE on STN AN85209798 MEDLINE DN PubMed ID: 2582083 Identification of Epstein-Barr virus-TΙ induced polypeptides in P3HR-1 cells by protein immunoblot. ΑU Sculley T B; Sculley D G; Pope J H SO Journal of general virology, (1985 May) 66 (Pt 5) 1113-22. Journal code: 0077340. ISSN: 0022-1317. CY ENGLAND: United Kingdom DTJournal; Article; (JOURNAL ARTICLE) LΑ English FS Priority Journals EM198507 ED Entered STN: 19900320 Last Updated on STN: 19990129 Entered Medline: 19850705 The protein immunoblot technique was used to identify ΔR Epstein-Barr virus-specific antigens present in sodium butyrate-induced P3HR-1 cells. Using sera from patients with either nasopharyngeal carcinoma or arthritis, 16 polypeptides were detected ranging in molecular weight from 22K to 140K. Each of the anti-EA-, anti-VCA-positive sera were found to contain antibodies to different subsets of the antigens. A 72K protein was identified which was consistent with the nuclear antigen (EBNA), and culturing cells in the presence of disodium phosphonoacetate allowed identification of 140K and 22K antigens as late viral products. Treatment of cells with sodium butyrate revealed that expression of some antigens

increased in parallel with the time of incubation of the cells in butyrate while other antigens either appeared early and then decreased in intensity or were only present after a number of days of butyrate treatment. One of

the antigens which decreased with the time cells were treated with

butyrate was EBNA.

```
L11 ANSWER 11 OF 47
                         MEDLINE on STN
AN
     84263215
                  MEDLINE
DN
     PubMed ID: 6086504
     Variation in expression of mouse erythrocyte receptors on
TI
     Epstein-Barr virus-induced B-cell
     lines.
ΑU
     Youinou P Y; Walker P R; Irving W L; Lydyard P M
SO
     Immunology letters, (1984) 8 (1) 27-32.
     Journal code: 7910006. ISSN: 0165-2478.
CY
     Netherlands
DT
     Journal; Article; (JOURNAL ARTICLE)
LΑ
     English
FS
     Priority Journals
EM
     198409
ED
     Entered STN: 19900320
     Last Updated on STN: 19900320
     Entered Medline: 19840917
AB
     To explain the variation in the percentage of mouse erythrocyte
     rosette-forming cells (MERFC) during culture of Epstein-
     Barr virus (EBV) - induced B-cell
     lines, we provide evidence that (i) there is an altered expression of
     mouse red blood cell (MRBC) receptors on cell line cells during
     the mitotic cycle, and (ii) putative receptor-negative cells are
     capable of de novo synthesis of the receptor, and passively
     adsorbing receptor shed from receptor-positive cells.
L11 ANSWER 12 OF 47
                         MEDLINE on STN
ΑÑ
     84174062
                 MEDLINE
DN
     PubMed ID: 6324457
ΤI
     Production of monoclonal antibody to a late intracellular Epstein
     -Barr virus-induced antigen.
ΑU
     Kishishita M; Luka J; Vroman B; Poduslo J F; Pearson G R
NC
     CA 20679 (NCI)
SO
     Virology, (1984 Mar) 133 (2) 363-75.
     Journal code: 0110674. ISSN: 0042-6822.
CY
    United States
DT
     Journal; Article; (JOURNAL ARTICLE)
T.A
     English
FS
     Priority Journals
EM
     198405
ED
     Entered STN: 19900319
    Last Updated on STN: 19980206
     Entered Medline: 19840518
AΒ
    A monoclonal antibody designated L2 was produced against a late
     intracellular protein induced by Epstein-
     Barr virus (EBV). This protein was expressed in cells
    producing virus but not in EBV genome-positive nonproducer cell lines, EBV
    genome-negative cell lines, or producer cultures cultivated in the
    presence of phosphonoacetic acid as determined by immunofluorescence. In
    addition, the antibody did not react with the membranes of infected cells
     indicating that it was not directed against an EBV-induced membrane
    antigen component. The monoclonal antibody was shown to recognize a
    glycoprotein with a molecular weight of approximately 125K by
    SDS-polyacrylamide gel electrophoresis. This glycoprotein was
    consistently found to be slightly larger when isolated from the P3HR-1
    cell line as opposed to the B-95-8 cell line. A similar difference was
    also noted by comparison of peptide maps of this protein isolated by
     immunoaffinity chromatography from the two cell lines. Serological
```

studies indicated that this 125K glycoprotein was a major component of the

viral capsid-antigen (VCA) complex as defined by immunofluorescence.

```
15 OF 47
             MEDLINE on STN
AN
     83012388
                  MEDLINE
DN
     PubMed ID: 6289060
ΤI
     Epstein-Barr virus induced
     proteins V: comparison of EBV-specific polypeptides from
     different virus strains.
ΑU
     Georg-Fries B; Mueller-Lantzsch N
     Medical microbiology and immunology, (1982) 171 (1) 11-21.
SO
     Journal code: 0314524. ISSN: 0300-8584.
     GERMANY, WEST: Germany, Federal Republic of
CY
DT
     Journal; Article; (JOURNAL ARTICLE)
T.A
FS
     Priority Journals
EΜ
     198212
     Entered STN: 19900317
ED
     Last Updated on STN: 19980206
     Entered Medline: 19821202
     EBV-associated polypeptides induced in different Epstein-Barr Virus
AB
```

(EBV) -producing cell lines by the tumor promotor TPA, and from purified EBV particles derived from the same lines were radioactivity labeled and analyzed by immunoprecipitation with human VCA+MA+ sera. In virus-producing cells no significant differences in the molecular weight of 35S-methionine-labeled EBV-associated polypeptide patterns could be observed. The analysis 125I-labeled polypeptides from purified virus particles of four different strains revealed that, in addition to common polypeptides, individual EBV strains contain strain-specific high molecular weight glycopolypeptides. These polypeptides, constituting part of the membrane antigen complex, are present in varying amounts. While P3HR-1 virus particles contain a major component of 250 000 and small amounts of 340 000 molecular weight polypeptides, Q IMR-WIL virus particles have more 340 00 than 240 000 molecular weight polypeptides. Furthermore, in B95-8 particles and in particles from an EBV strain isolated from an African green monkey (AGM-EBV) respectively, large amounts of 360 000 and 250 000 polypeptides could be observed. Since these glycopolypeptides carry strain-, subgroup- and group-specific antigenic determinants, also found in virus strains produced in human and marmoset cells, it should be further investigated whether these differences in molecular weight are virus-strain- or cell-specific.

```
ANSWER 17 OF 47 MEDLINE on STN
L11
AN
     82054725
                  MEDLINE
DN
     PubMed ID: 6271909
TI
     The regulated expression of Epstein-Barr
     virus. III. Proteins specified by EBV
     during the lytic cycle.
ΑU
     Bayliss G J; Wolf H
SO
     Journal of general virology, (1981 Sep) 56 (Pt 1) 105-18.
     Journal code: 0077340. ISSN: 0022-1317.
CY
     ENGLAND: United Kingdom
DT
     Journal; Article; (JOURNAL ARTICLE)
LΑ
     English
FS
     Priority Journals
EM
     198201
ED
     Entered STN: 19900316
     Last Updated on STN: 19900316
     Entered Medline: 19820120
     The experiments show that 30 virus-induced or
AΒ
```

virus-specified proteins were synthesized in

Raji cells after superinfection with <code>Epstein-Barr virus</code> (EBV) derived from P3HR1 cells. Using a combination of pulse labelling, application of cycloheximide blocks at different times post-infection, treatment with amino acid analogues and inhibition of DNA synthesis it was shown that three groups of proteins appear in Raji cells after superinfection; the synthesis of the proteins in any one group appears to be coordinately regulated. Amongst the six <code>virus-induced proteins</code> which were synthesized immediately after release from an early cycloheximide block one would expect to find those <code>proteins</code> essential for the transition from EBNA to EA synthesis. Using human sera with differing specificities for the various antigen groups 11 <code>proteins</code> were identified as being specifically precipitated by sera having high titres against the <code>EBV-induced</code> early antigen complex.

```
L11 ANSWER 20 OF 47
                         MEDLINE on STN
ΑN
     80170704
                 MEDLINE
DN
     PubMed ID: 6154378
     Epstein-Barr virus-induced
     proteins. II. Analysis of surface polypeptides from EBV
     -producing and -superinfected cells by immunoprecipitation.
     Meuller-Lantzsch N; Georg B; Yamamoto N; zur Hausen H
ΑU
     Virology, (1980 Apr 30) 102 (2) 401-11.
SO
     Journal code: 0110674. ISSN: 0042-6822.
CY
     United States
DT
     Journal; Article; (JOURNAL ARTICLE)
LΑ
     English
FS
     Priority Journals
ΕM
     198006
ED
     Entered STN: 19900315
     Last Updated on STN: 19900315
     Entered Medline: 19800625
L11 ANSWER 21 OF 47
                         MEDLINE on STN
     80170682
AN
                 MEDLINE
DN
     PubMed ID: 6245505
TI
     Epstein-Barr virus-induced
     proteins. III. Analysis of polypeptides from P3HR-1-EBV
     -superinfected NC37 cells by immunoprecipitation.
     Mueller-Lantzsch N; Georg B; Yamamoto N; zur Hausen H
ΑU
SO
     Virology, (1980 Apr 15) 102 (1) 231-3.
     Journal code: 0110674. ISSN: 0042-6822.
CY
     United States
DT
     Journal; Article; (JOURNAL ARTICLE)
ĽΑ
     English
FS
     Priority Journals
EM
     198006
     Entered STN: 19900315
ED
     Last Updated on STN: 19900315
     Entered Medline: 19800616
L11 ANSWER 22 OF 47
                       MEDLINE on STN
AN
     79238644 MEDLINE
DN
     PubMed ID: 89099
```

membrane antigens: immunochemical characterization of Triton X-100 solubilized viral membrane antigens from EBV-superinfected Raji

International journal of cancer. Journal international du cancer,

ΤI

ΑU

SO

Epstein-Barr virus-induced

Qualtiere L F; Pearson G R

(1979 Jun 15) 23 (6) 808-17.

Journal code: 0042124. ISSN: 0020-7136.

CY Denmark

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 197910

> Last Updated on STN: 19900315 Entered Medline: 19791017

AB In an attempt to qualitatively identify the membrane antigen (MA) complex induced by Epstein-Barr virus (EBV)

infection of lymphoblastoid cells, superinfected Raji cells were surface labelled with 125I by the lactoperoxidase method and solubilized with Triton X-100, then the 125I-labelled membrane proteins were precipitated by sera containing high antibody titers to MA. Analysis of these immune precipitates on sodium dodecyl sulfate polyacrylamide gel eletrophoresis identified four major EBV-specific membrane proteins with molecular weights (mol. wt) of 280,000, 250,000, 170,000 and 90,000. Sera from patients with Burkitt's lymphoma (BL), nasopharyngeal carcinoma (NPC) and infectious mononucleosis (IM) and from EBV-infected disease-free individuals showed differential patterns of reactivity to these antigens with some sera only recognizing three or less of the antigens. The results from invesigations with these sera also indicated that these major proteins were not related to EBV-induced viral

capsid antigens (VCA) or the virus-associated early antigen (EA) complexes as defined by immunofluorescence. Metabolic labelling of EBV-infected Raji cells with [14C]glucosamine, followed by Triton X-100 solubilization and radioimmune precipitation, identified the 280,000, 250,000 and 90,000 components as glycoproteins. The lactoperoxidase-labelled 170,000 molecular weight component was not significantly glycosylated and, therefore, could not be categorized as a glycoprotein on the basis of this study. In addition, a glycoprotein with a mol. wt of 130,000 was identified by this approach which also appeared to be specified by EBV. The results from these investigations, therefore, indicated that the EBV-induced MA complex was composed of four major glycoproteins and one nonglycosylated high mol. wt protein.

```
L11 ANSWER 24 OF 47 CAPLUS COPYRIGHT 2005 ACS on STN
```

AN 1994:104348 CAPLUS

DN 120:104348

TI Analysis of human T cell responses to Epstein-Barr virus-induced replication proteins

AU Varghese, Susan

CS Med. Cent., Georgetown Univ., Washington, DC, USA

SO (1992) 281 pp. Avail.: Univ. Microfilms Int., Order No. DA9236641

From: Diss. Abstr. Int. B 1993, 53(7), 3394

DT Dissertation

LA English

AB Unavailable

- L11 ANSWER 26 OF 47 CAPLUS COPYRIGHT 2005 ACS on STN
- AN 1990:438326 CAPLUS
- DN 113:38326
- TI Protein kinase activity associated with an Epstein-Barr virus-induced early antigen
- AU Kocache, Malda Mahmoud
- CS Georgetown Univ., Washington, DC, USA

SO (1989) 189 pp. Avail.: Univ. Microfilms Int., Order No. DA9006752

From: Diss. Abstr. Int. B 1990, 50(10), 4367

DT Dissertation

LA English

AB Unavailable

- L11 ANSWER 27 OF 47 CAPLUS COPYRIGHT 2005 ACS on STN
- AN 1985:40774 CAPLUS
- DN 102:40774
- TI Mapping of genes involved in Epstein-Barr virus (EBV)-induced lymphocyte transformation
- AU Volsky, D. J.; Gross, T.; Volsky, B.; Bartzatt, R.; Kuszynski, C.; Sinangil, F.
- CS Med. Cent., Univ. Nebraska, Omaha, NE, 68105, USA
- SO UCLA Symposia on Molecular and Cellular Biology, New Series (1984), 17(Genes Cancer), 293-302
 CODEN: USMBD6; ISSN: 0735-9543
- DT Journal
- LA English
- AB Purified EBV DNA (B-95-8 strain) and cloned DNA fragments were trapped in Sendai virus envelopes during envelope reconstitution. The fusogenic DNA-loaded envelopes (RSVE/DNA) served as gene transfer vehicles for mapping the EBV genome in fresh human lymphocytes (HL). EBV DNA induced EBV-determined nuclear antigen (EBNA) in 0.2-1% of HL, transiently stimulated cellular DNA synthesis, but did not fully transform cells. HL were transformed after coinfection with viral DNA and UV-inactivated B-95-8 EBV. Cloned SalI F1 (9 kilobase pairs (kbp)) and a smaller BamHI K (5.2 kbp) fragment from the same region of EBV DNA induced EBNA in 0.2-4% of HL but did not stimulate cellular DNA synthesis nor transform cells. Cloned BamHI D1 fragment (9 kbp) from AG876 virus DNA stimulated cellular DNA synthesis but did not induce EBNA. EA and VCA were not observed with any of the DNA fragments tested. Apparently, induction of EBNA alone is not sufficient for achieving transformation of human lymphocytes.
- L11 ANSWER 44 OF 47 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- AN 81:535220 SCISEARCH
- GA The Genuine Article (R) Number: MP021
- TI EPSTEIN-BARR VIRUS-INDUCED
 PROTEINS 4. CHARACTERIZATION OF AN EBV-ASSOCIATED
 PHOSPHOPOLYPEPTIDE
- AU MUELLERLANTZSCH N (Reprint); YAMAMOTO N
- CS UNIV FREIBURG, INST VIROL, ZENTRUM HYG, D-7800 FREIBURG, FED REP GER
- CYA FEDERAL REPUBLIC OF GERMANY
- SO JOURNAL OF GENERAL VIROLOGY, (1981) Vol. 55, No. AUG, pp. 333-341.
- DT Article; Journal
- FS LIFE
- LA ENGLISH
- REC Reference Count: 29
- L11 ANSWER 45 OF 47 LIFESCI COPYRIGHT 2005 CSA on STN
- AN 87:17037 LIFESCI
- TI Characterization of the Epstein-Barr virusinduced early polypeptide complex p50/58 EA-D using rabbit antisera, a monoclonal antibody, and human antibodies.
- AU Doelken, G.; Hecht, T.; Roeckel, D.; Hirsch, F.W.
- CS City of Hope Natl. Med. Cent., Dep. Hematol. and Bone Marrow Transplant., 1500 E. Duarte Rd., Duarte, CA 91010, USA

- SO VIROLOGY., (1987) vol. 157, no. 2, pp. 460-471.
- DT Journal
- FS V
- LA English
- SL English
- AB A polypeptide complex (p52) belonging to the D-subspecificity of the EBV-induced early antigens (EA-D) was purified from chemically induced P3HR-1 cells. Rabbit antisera raised against the isolated polypeptides reacted with components of EA-D as could be shown by indirect immunofluorescence and immunoperoxidase staining of IdU-induced EA positive Raji cells, ELISA, and immunoblotting. In one-dimensional immunoblots the rabbit antisera detected a predominant polypeptide complex of 52 kDa. Two-dimensional immunoblots prepared with proteins from IdU-induced Raji cells showed that the rabbit sera detect three series of polypeptides of 52 kDa (p/8.5-6.2), 55-58 kDa (p/6.2-4.5), and 48-50 kDa (p/6.0-4.5). These three groups of polypeptides could also be identified by 50 high titered anti-EA-D positive human sera and a specific monoclonal antibody (R3) as being the main components of EA-D in Raji and B95-8 cells.
- L11 ANSWER 46 OF 47 LIFESCI COPYRIGHT 2005 CSA on STN
- AN 83:77290 LIFESCI
- TI Purification of a protein (60K/58K) associated with the **Epstein-Barr virus-induced** early antigen complex in Raji cells.
- AU Doelken, G.; Lange, W.; Weitzmann, U.; Hirsch, F.W.; Loehr, G.W.
- CS Abt. Haematol. und Onkol., Med. Universitaetsklin., Freiburg, Hugstetter-Str. 55, D-7800 Freiburg, FRG
- SO INT. J. CANCER., (1983) vol. 32, no. 3, pp. 307-314.
- DT Journal
- FS V; F; L
- LA English
- SL English
- AB A double antibody sandwich ELISA has been established for the detection and quantification of EBV-associated early antigens (EA) in IUdR-induced Raji cells. The EA complex extracted from Raji cells could be separated by ion exchange chromatography and isoelectric focusing into several components. One EA-associated subspecificity has been purified by DEAE-, CM-, and Blue-Sepharose chromatography followed by isoelectric focusing. The isolated protein has an apparent molecuar weight of 240,000 plus or minus 20,000 daltons under non-dissociating conditions on Sepharcyl S-300, an isoelectric point of 4.5, and seems to be composed of two polypeptides of 60,000 and 58,000 daltons as shown by SDS-gel electrophoresis and two-dimensional gel electrophoresis. The EA activity of the isolated protein has been confirmed by the double antibody sandwich ELISA and its reactivity with anti-EA-positive sera in an ELISA for the detection of anti-EA antibodies.

WEST Search History

Hide Items Restore Clear Cancel

DATE: Tuesday, March 01, 2005

Hide?	<u>Set</u> Name	Query	<u>Hit</u> Count
DB=PGPB,USPT,EPAB,JPAB,DWPI; PLUR=YES; OP=OR			
	L21	120 and 112	42
	L20	19921125	108
	L19	19921125	1
	L18	L13 and (binding adj fragment or binding-fragment) same antibod\$4	1137
	L17	L14 and (binding adj fragment or binding-fragment) same antibod\$4	1
	L16	19921125	2
	L15	113 and (affinity adj label) same (avidin or biotin)	353
	L14	19921125	9
	L13	111 and L12	2628
	L12	(antibod\$3 or immunoglob\$4) same label and label with (peroxidase or biotin or rhodamine)	15241
	L11	(antibod\$3 or immunoglob\$4) same label and (peroxidase same biotin same rhodamine)	3446
	L10	(antibod\$3 or immunoglob\$4) same label and (peroxidase or biotin or rhodamine)	24929
	L9	(epstein adj barr adj3 \$induced or epstein-barr-induced)	208
	L8	L7 not 14	8
	L7	L6 and l1	26
	L6	(epstein adj barr adj3 induced or epstein-barr-induced)	208
	L5	(epstein adj barr adj induced or epstein-barr-induced)	5
	L4	((lymphoid-specific or lymphoid adj specific) with (G-protein-coupled or (G-protein or g adj protein) adj coupled) adj receptor) or (EBI1 or EBI-1) and epstein adj barr or epstein adj barr adj induced or epstein-barr-induced	43
	L3	1995	23
	L2	L1 and (antibod\$ or immunoglob\$)	791
	L1	(lymphoid-adj specific with (G-protein-coupled or (G-protein or g adj protein) adj coupled) adj receptor) or (EBI1 or EBI-1) and epstein adj barr or epstein adj barr adj induced or epstein-barr-induced	981

END OF SEARCH HISTORY